

4164-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2011-N-0781]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Record Retention Requirements for the Soy Protein and Risk of Coronary Heart Disease Health Claim

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (PRA).

DATES: Fax written comments on the collection of information by [INSERT DATE 30 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER].

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, Fax: 202-395-7285, or emailed to oira_submission@omb.eop.gov. All comments should be identified with the OMB control number 0910-0428. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: JonnaLynn Capezzuto, Office of Operations, Food and Drug Administration, Three White Flint North, 10A-12M, 11601 Landsdown St., North Bethesda, MD 20852, 301-796-3794, PRAStaff@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Record Retention Requirements for the Soy Protein and Risk of Coronary Heart Disease Health

Claim--21 CFR 101.82

OMB Control Number 0910-0428--Extension

Section 403(r)(3)(A) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 343(r)(3)(A)) provides for the use of food label statements characterizing a relationship of any nutrient of the type required to be in the label or labeling of the food to a disease or a healthrelated condition only where that statement meets the requirements of the regulations issued by the Secretary of Health and Human Services to authorize the use of such a health claim. Section 101.82 (21 CFR 101.82) of our regulations authorizes a health claim for food labels about soy protein and the risk of coronary heart disease. Accordingly, FDA established the previously referenced information collection in support of the regulation. In the Federal Register of October 31, 2017 (82 FR 50324), we published a proposed rule to revoke the underlying regulation found at § 101.82. We are taking this action based on our review of the totality of publicly available scientific evidence currently available and our tentative conclusion that such evidence does not support our previous determination that there is significant scientific agreement among qualified experts for a health claim regarding the relationship between soy protein and reduced risk of coronary heart disease. Upon finalization of the proposed rule, the associated information collection requirements under this OMB control number will be revoked. 3

Until such time and in accordance with the PRA, we retain our currently approved burden

estimate for the information collection displayed in table 1 of this notice.

In the Federal Register of March 8, 2018 (83 FR 9856), FDA published a 60-day notice

requesting public comment on the proposed collection of information. No comments were

received.

FDA estimates the burden of this collection of information as follows:

Table 1.--Estimated Annual Recordkeeping Burden¹

	1 6				
21 CFR Section	No. of	No. of Records per	Total	Average Burden	Total
	Recordkeepers	Recordkeeper	Annual	per	Hours
			Records	Recordkeeping	
101.82(c)(2)(ii)(B)	25	1	25	1	25

There are no capital costs or operating and maintenance costs associated with this collection of information.

Based on our current experience with the use of health claims, we estimate 25 firms

market products bearing a soy protein/coronary heart disease health claim and that perhaps one

of each firm's products might contain non-soy sources of protein along with soy protein. The

records currently required to be retained under § 101.82(c)(2)(ii)(B) are the records, e.g., the

formulation or recipe, that a manufacturer has and maintains as a normal course of its doing

business. Thus, the burden to the food manufacturer is limited to assembling and retaining the

records, which we estimate will take 1 hour annually.

Dated: May 30, 2018.

Leslie Kux,

Associate Commissioner for Policy.

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